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In the claims:

Please amend claims as follows:

1-20. (Cancelled)

21. (Currently amended) A non-human transgenic <u>animal organism</u> having a transgene integrated into the genome of the <u>animal organism</u> and also having a *tet* operator-linked gene in the genome of the <u>animal organism</u>, wherein:

the transgene comprises a transcriptional regulatory element functional in cells of the <u>animal organism</u> operatively linked to a polynucleotide sequence encoding a fusion protein which activates transcription of said *tet* operator linked gene,

the fusion protein comprises a first polypeptide which is a mutated Tet repressor that binds to a *tet* operator sequence in the presence of tetracycline or a tetracycline analogue operatively linked to a second polypeptide which activates transcription in eukaryotic cells,

said tet operator linked gene confers a detectable and functional phenotype on the organism when expressed in cells of the animal,

said transgene is expressed in cells of the animal at a level sufficient to produce amounts of said fusion protein that are sufficient to activate transcription of the *tet* operator-linked gene at detectable levels; and

in the presence of tetracycline or a tetracycline analogue in the <u>animal organism</u>, said fusion protein binds to the *tet* operator-linked gene and activates transcription of the *tet* operator linked gene such that the *tet* operator-linked gene is expressed at <u>detectable</u> <u>levels a level sufficient to confer the detectable and functional phenotype on the animal</u>, wherein the level of expression of the *tet* operator-linked gene can be downmodulated by depleting tetracycline or a tetracycline analogue from the animal.

22. (Currently amended) A non-human transgenic <u>animal organism</u> having a transgene integrated into the genome of the <u>animal organism</u> and also having a *tet* operator-linked gene in the genome of the <u>animal organism</u>, wherein:

the transgene comprises a transcriptional regulatory element functional in cells of the <u>animal organism</u> operatively linked to a polynucleotide sequence encoding a fusion protein which inhibits transcription of said *tet* operator linked gene,

the fusion protein comprises a first polypeptide which is a mutated Tet repressor that binds to a *tet* operator sequence in the presence of tetracycline or a tetracycline

analogue operatively linked to a second polypeptide which inhibits transcription in eukaryotic cells,

said tet operator-linked gene confers a detectable and functional phenotype on the organism when expressed in cells of the organism,

said transgene is expressed in cells of the <u>animal organism</u> at a level sufficient to produce amounts of said fusion protein that are sufficient to inhibit transcription of the *tet* operator-linked gene; and

in the presence of tetracycline or a tetracycline analogue in the <u>animal organism</u>, said fusion protein binds to the *tet* operator-linked gene and inhibits transcription of the *tet* operator linked gene, wherein the level of expression of the *tet* operator-linked gene can be upregulated by depleting tetracycline or a tetracycline analogue from the <u>animal organism</u>.

23. (Currently amended) A non-human transgenic <u>animal organism</u> having a transgene integrated into the genome of the <u>animal organism</u> and also having a *tet* operator-linked gene in the genome of the <u>animal organism</u>, wherein:

the transgene comprises a transcriptional regulatory element functional in cells of the <u>animal organism</u> operatively linked to a polynucleotide sequence encoding a fusion protein which inhibits transcription of said *tet* operator linked gene,

said fusion protein comprises a first polypeptide that is a Tet repressor, operably linked to a heterologous second polypeptide which inhibits transcription of said *tet* operator-linked gene in eucaryotic cells,

said tet operator linked gene confers a detectable and functional phenotype on the organism when expressed in cells of the organism,

said transgene is expressed in cells of the <u>animal organism</u> at a level sufficient to produce amounts of said fusion protein that are sufficient to inhibit transcription of the *tet* operator-linked gene; and

in the absence of tetracycline or a tetracycline analogue in the organism, said fusion protein binds to the *tet* operator-linked gene and inhibits transcription of the *tet* operator linked gene, wherein the level of expression of the *tet* operator-linked gene can be upregulated by administering tetracycline or a tetracycline analogue to the <u>animal</u> organism.

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24. (Currently amended) The <u>animal organism</u> of claim 21, wherein the mutated Tet repressor has at least one amino acid substitution compared to a wild-type Tet repressor.

- 25. (Currently amended) The <u>animal organism</u> of claim 24, wherein the mutated Tet repressor is a mutated Tn10- derived Tet repressor having an amino acid substitution at at least one amino acid position selected from the group consisting of position 71, position 95, position 101 and position 102.
- 26. (Currently amended) The <u>animal organism</u> of claim 24, wherein the mutated Tn10-derived Tet repressor comprises an amino acid sequence shown in positions 1 to 207 of SEQ ID NO: 2.
- 27. (Currently amended) The <u>animal organism</u> of claim 21, wherein the second polypeptide of the fusion protein comprises a transcription activation domain of herpes simplex virion protein 16.
- 28. (Currently amended) The <u>animal organism</u> of claim 22, wherein the second polypeptide of the fusion protein comprises a transcriptional silencer domain of a protein selected from the group consisting of v-erbA, the Drosophila Krueppel protein, the retinoic acid receptor alpha, the thyroid hormone receptor alpha, the yeast Ssn6/Tup1 protein complex, the Drosophila protein even-skipped, SIR1, NeP1, the Drosophila dorsal protein, TSF3, SF1, the Drosophila hunchback protein, the Drosophila knirps protein, WT1, Oct-2.1, the Drosophila engrailed protein, E4BP4 and ZF5.
- 29. (Currently amended) The <u>animal organism</u> of claim 21, wherein expression of the transgene is regulated by at least one *tet* operator sequence.
- 30. (Currently amended) The <u>animal organism</u> of claim 21, wherein expression of the transgene is regulated by at least one virally-derived regulatory element.
- 31. (Currently amended) The <u>animal organism</u> of claim 21, wherein expression of the transgene is regulated by at least one tissue-specific regulatory element.

32. (Currently amended) The <u>animal organism</u> of claim 21, wherein the *tet* operator-linked gene is a second transgene comprising a gene of interest operably linked to at least one *tet* operator sequence.

- 33. (Currently amended) The <u>animal organism</u> of claim 32, wherein the at least one *tet* operator sequence is operatively linked upstream of the second transgene.
- 34. (Currently amended) The <u>animal organism</u> of claim 32, wherein the at least one *tet* operator sequence is operatively linked downstream of the second transgene.
- 35. (Currently amended) The <u>animal organism</u> of claim 21, wherein the *tet* operator-linked gene is an endogenous gene that has been operatively linked to at least one *tet* operator sequence.
- 36. (Currently amended) The <u>animal organism</u> of claim 21, which is selected from the group consisting of: <u>a mouse</u>, <u>a goat</u>, a cow, a sheep, <u>a monkey</u>, a dog, a cat, a rabbit, a rat, or a pig.
- 37. (New) A transgenic non-human animal whose genome comprises a transgene comprising a transcriptional regulatory element functional in cells of the transgenic non-human animal operatively linked to a polynucleotide sequence encoding a fusion protein,

said fusion protein comprising a first polypeptide which is a Tet repressor operatively linked to a second polypeptide which directly or indirectly activates transcription of a *tet* operator linked gene such that the transgenic non-human animal has detectable levels of the *tet* operator linked gene in one or more cell types or tissues of the transgenic non-human animal.

- 38. (New) The transgenic non-human animal of claim 37, wherein the Tet repressor is a mutant Tet repressor.
- 39. (New) The transgenic non-human animal of claim 38, wherein the mutated Tet repressor has at least one amino acid substitution compared to a wild-type Tet repressor.

40. (New) The transgenic non-human animal of claim 38, wherein the mutated Tet repressor is a mutated Tn10- derived Tet repressor having an amino acid substitution at at least one amino acid position selected from the group consisting of position 71, position 95, position 101 and position 102.

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- 41. (New) The transgenic non-human animal of claim 38, wherein the mutated Tn10-derived Tet repressor comprises an amino acid sequence shown in positions 1 to 207 of SEQ ID NO: 2.
- 42. (New) The transgenic non-human animal of claim 37, wherein the second polypeptide of the fusion protein comprises a transcription activation domain of herpes simplex virion protein 16.
- 43. (New) The animal of claim 37, which is selected from the group consisting of a mouse, a goat, a cow, a sheep, a monkey, a dog, a cat, a rabbit, a rat, or a pig.
- 44. (New) A transgenic non-human animal whose genome comprises a *tet* operator-linked gene and

a transgene comprising a transcriptional regulatory element functional in cells of the transgenic non-human animal operatively linked to a polynucleotide sequence encoding a fusion protein which activates transcription of the *tet* operator linked gene,

said fusion protein comprising a first polypeptide which is a Tet repressor operatively linked to a second polypeptide which directly or indirectly activates transcription in eukaryotic cells,

wherein said transgene is expressed in cells of the transgenic non-human animal at a level sufficient to produce amounts of said fusion protein that are sufficient to activate transcription of the *tet* operator-linked gene at detectable levels in the absence of tetracycline or a tetracycline analogue.

45. (New) A transgenic non-human animal whose genome comprises a *tet* operator-linked gene and

a transgene comprising a transcriptional regulatory element functional in cells of the transgenic non-human animal operatively linked to a polynucleotide sequence encoding a fusion protein which activates transcription of the *tet* operator linked gene, said fusion protein comprising a first polypeptide which is a mutated Tet repressor operatively linked to a second polypeptide which directly or indirectly activates transcription in eukaryotic cells,

wherein said transgene is expressed in cells of the transgenic non-human animal at a level sufficient to produce amounts of said fusion protein that are sufficient to activate transcription of the *tet* operator-linked gene at detectable levels in the presence of tetracycline or a tetracycline analogue.

- 46. (New) The transgenic non-human animal of claim 45, wherein the mutated Tet repressor has at least one amino acid substitution compared to a wild-type Tet repressor.
- 47. (New) The transgenic non-human animal of claim 46, wherein the mutated Tet repressor is a mutated Tn10- derived Tet repressor having an amino acid substitution at at least one amino acid position selected from the group consisting of position 71, position 95, position 101 and position 102.
- 48. (New) The transgenic non-human animal of claim 46, wherein the mutated Tn10-derived Tet repressor comprises an amino acid sequence shown in positions 1 to 207 of SEQ ID NO: 2.
- 49. (New) The transgenic non-human animal of claim 44, wherein the second polypeptide of the fusion protein comprises a transcription activation domain of herpes simplex virion protein 16.
- 50. (New) The transgenic non-human animal of claim 45, wherein the second polypeptide of the fusion protein comprises a transcription activation domain of herpes simplex virion protein 16.
- 51. (New) A transgenic non-human animal whose genome comprises a *tet* operator-linked gene and
- a transgene comprising a transcriptional regulatory element functional in cells of the transgenic non-human animal operatively linked to a polynucleotide sequence encoding a fusion protein which inhibits transcription of the *tet* operator linked gene,

said fusion protein comprising a first polypeptide which is a Tet repressor operatively linked to a second polypeptide which directly or indirectly inhibits transcription in eukaryotic cells,

wherein said transgene is expressed in cells of the transgenic non-human animal at a level sufficient to produce amounts of said fusion protein that are sufficient to inhibit transcription of the *tet* operator-linked gene.

- 52. (New) The transgenic non-human animal of claim 51, wherein the second polypeptide of the fusion protein comprises a transcriptional silencer domain of a protein selected from the group consisting of v-erbA, the Drosophila Krueppel protein, the retinoic acid receptor alpha, the thyroid hormone receptor alpha, the yeast Ssn6/Tup1 protein complex, the Drosophila protein even-skipped, SIR1, NeP1, the Drosophila dorsal protein, TSF3, SF1, the Drosophila hunchback protein, the Drosophila knirps protein, WT1, Oct-2.1, the Drosophila engrailed protein, E4BP4 and ZF5.
- 53. (New) The transgenic non-human animal of claim 51, wherein the Tet repressor is a mutant Tet repressor.
- 54. (New) The animal of claim 51, which is selected from the group consisting of a mouse, a goat, a cow, a sheep, a monkey, a dog, a cat, a rabbit, a rat, or a pig.